

## **SUPPLEMENTARY MATERIAL 1**

### **PCR amplification of rs13337493 (restriction fragment length polymorphism analysis)**

In the polymerase chain reaction (PCR) amplification, 1  $\mu\text{L}$  of genomic DNA (10  $\text{ng}/\mu\text{L}$ ) was applied to multiplex PCR reaction in 20  $\mu\text{L}$  containing 0.5  $\mu\text{M}$  of each primer and 10  $\mu\text{L}$  of 2 $\times$  Taq-Plus Master Mix (ACE Biosystems, Taoyuan, Taiwan). The following thermal cycling procedure was employed: 95°C for 5 min; 40 cycles of 95°C for 40 s, 52°C for 30 s, and 72°C for 40 s; extension at 72°C for 10 min; and finally, a hold cycle at 4°C.